CLEAN VERSION OF AMENDMENTS

IN THE CLAIMS

Please amend claims 1 and 3 to read as follows:

1. (amended) A process for producing solid dosage forms which are suitable for oral or rectal administration for humans and animals, wherein

a) 0.5 to 30% by weight of at least one active ingredient,

b) 0.5 to 70% by weight of at least one cyclodextrin,

c) 10 to 98% by weight of at least one polymeric binder, selected from the group consisting of polyethylene glycol having a molecular weight above 1000, polyvinylpyrrolidone, and copolymers comprising N-vinylpyrrolidone and vinyl acetate, and

d) 0 to 50% by weight of conventional excipients

are mixed and plasticized at a temperature below 220°C without adding a solvent and the resulting plastic mixture is shaped to produce the dosage form.

3. (twice amended) A process as claimed in claim 1, wherein the plastic mixture is shaped in a molding calendar to produce the dosage forms.